## IN THE CLAIMS:

Please enter the attached listing of claims into the application. This listing of claims replaces all prior listing of claims in the application.

## LISTING OF CLAIMS

- 1-48. (Cancelled)
- 49. (Currently Amended) A method for assessing the risk of colorectal cancer and colorectal polyps, comprising:
- selecting a panel of biomarkers comprising at least two polynucleotides including SEQ ID Nos:1 and 2;

amplifying and quantifying RNA expression levels in a biological colorectal sample <u>obtained from normal appearing mucosa</u> from a subject for each biomarker in the panel including polynucleotides comprising SEQ ID Nos:1 and 2; <del>and</del>

comparing the quantified expression levels of each biomarker including polynucleotides comprising SEQ ID NO:1 and 2 in the sample to each of the same biomarker expression level in a normal control colorectal sample; and

determining a difference in the expression levels in the biological sample compared to the normal control wherein an increase in the expression level of at least SEQ ID NO:1 and/or 2 is indicative of an increased risk of colorectal cancer and colorectal polyps.

- 50. (Previously Presented) The method of claim 49, where the step of selecting a panel of biomarkers further comprises at least one polynucleotide having a sequence selected from the group consisting of SEQ ID NOs:3-15 and 16.
- 51. (Previously Presented) The method of claim 49, where the step of selecting a panel of biomarkers further comprises at least one polynucleotide comprising a sequence selected from SEQ ID Nos: 15 and 16.
- (Previously Presented) The method of claim 51, where the step of amplifying further comprises using at least two sets of primers chosen from (i) SEQ

ID NO:45 and 46, (ii) SEQ ID NO:47 and 48, (iii) SEQ ID NO:53 and 54, (iv) SEQ ID NO:73 and 74 and (v) SEQ ID NO:75 and 76.

- (Previously Presented) The method of claim 52, where the step of amplifying further comprises using enzymes and reagents for the preparation of cDNAs.
- 54. (Previously Presented) The method of claim 49, where the step of quantifying the levels of RNA further comprises labeling the amplified polynucleotide.
- (Previously Presented) The method of claim 54, where labeling includes at least one chromophore.
- (Cancelled).
- 57. (Previously Presented) The method of claim 49, wherein an increase in a polynucleotide comprising SEQ ID NO:1 and/or 2 in the sample compared to levels of corresponding biomarkers from the normal control identifies the subject as a candidate for the risk management of colorectal cancer and colorectal polyps.
- 58-60. (Cancelled)
- 61. (Previously Presented) The method of claim 49, further comprising obtaining a sample of colorectal cells by minimally invasive or non-invasive techniques.
- 62. (Original) The method of claim 61, where the minimally invasive step is by use of a swab.
- 63. (Previously Presented) The method of claim 61, where obtaining a sample of colorectal cells is non-invasive

64-95. (Cancelled)

- 96. (Previously Presented) The method of claim 57, wherein the biomarker comprises (i) SEQ ID NO:1, (ii) SEQ ID NO:2, or (iii) SEQ ID Nos:1 and 2.
- 97. (Cancelled).
- 98. (Currently Amended) A method for assessing the risk of colorectal cancer, comprising:

selecting a panel of biomarkers comprising polynucleotides having sequences including SEQ ID Nos:1 and 2;

obtaining a biological colorectal sample <u>from normal appearing mucosa</u> from a subject:

isolating cellular RNA from the sample;

amplifying and quantifying RNA expression levels in a biological colorectal sample from a subject for each biomarker in the panel including SEQ ID Nos:1 and 2: and

comparing the quantified expression levels of each biomarker in the sample to each of the same biomarker expression level in a normal control colorectal sample; and

determining a difference in the expression levels of the biomarkers in the panel including SEQ ID Nos: 1 and 2 in the biological sample compared to the normal control, wherein an increase in at least SEQ ID NO:1 and/or 2 is indicative of an increased risk of colorectal cancer.

- (Previously Presented) The method of claim 98, where the step of selecting a panel of biomarkers further comprises at least one additional polynucleotide from SEQ ID NOs: 3-16.
- 100. (Previously Presented) The method of claim 98, where the step of quantifying the levels of RNA further comprises labeling cDNA.

- 101. (Previously Presented) The method of claim 100, where labeling cDNA includes at least one chromophore.
- 102. (Previously Presented) The method of claim 98, wherein an increase in the expression of a polynucleotide comprising SEQ ID NO:1 and/or 2 in the sample compared to levels of corresponding biomarkers from the normal control identifies the subject as a candidate for further clinical management.
- 103. (Previously Presented) The method of claim 98, where the step of obtaining a sample of colorectal cells is minimally invasive or non-invasive.
- 104. (Previously Presented) The method of claim 103, where the minimally invasive step is by use of a swab.
- 105. (Previously Presented) The method of claim 102, where the step of obtaining a sample of colorectal cells is non-invasive.
- 106-107. (Cancelled)